Dercum’s Disease: A Rare Disease of Painful Fatty Lumps

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Abstract

Dercum’s disease is a rare and poorly understood condition characterized by painful subcutaneous adipose tissue growth that can occur anywhere beneath the skin surface. We present the case of a 27-year-old man with no significant medical history who had been experiencing painful subcutaneous nodules for 2 years. Skin biopsy revealed proliferation of mature adipocytes that surrounded with fibrous septa. There are currently no treatments approved by the US Food and Drug Administration for Durum’s disease, and the effectiveness of treatments that have been attempted is variable.

Introduction

Adiposis dolorosa, also known as Dercum’s disease, is a rare and not properly understood condition characterized by painful subcutaneous adipose tissue growth that can occur anywhere beneath the skin’s surface [1]. Francis Dercum, an American neurologist, described it for the first time in 1888 [2]. Many related symptoms, including fatigue, depression, dementia, and weakness, are frequently present along with the disease [1]. The disease is inherited on an autosomal dominant basis with variable penetrance, and most of the cases are sporadic [3]. The pathophysiology remains unclear [4]. The epidemiology of this condition has not yet been defined, but literature suggests a higher prevalence in postmenopausal obese females [5]. We report a case of Dercum’s disease in a healthy male patient.

Case Presentation

We present the case of a 27-year-old man with no significant medical history who had been experiencing painful subcutaneous nodules for two years all over his body except his head, neck, hands, and feet. The condition started with a gradual onset and was progressive. He has developed an alarming rise in the quantity of these masses. The patient also experienced paresthesia. His quality of life was significantly reduced because of painful symptoms that frequently interrupted his daily activities. Pain was refractory to analgesics and anti-inflammatory drugs. There was no history of mental disorders. The systemic review was not significant. There was no family history of obesity. He was not an alcoholic or smoker.

On examination, there were multiple soft-tissue swellings, more in the trunk and upper limbs than lower limbs with sparing of the face, hands, and feet (figure 1). The swellings were painless to palpation. The patient’s BMI was 24 kg/m2. Neurological evaluation was normal, and other physical findings were unremarkable.

Laboratory investigations revealed an erythrocyte sedimentation rate of 8 mm/h and serum C-reactive protein level of 4 mg/L. Other laboratory tests such as blood cell counts, serum electrolytes, liver function tests, serum protein electrophoresis, and lipid profiles were normal. Skin biopsy revealed dermal mass formed of proliferating adipocytes surround with...
FIGURE 1: Multiple swellings scattered over the trunk, upper, and lower limbs.

FIGURE 2: Histopathological findings of Dercum’s disease

A) Low-power 10X, H&E-stained biopsy of the subcutaneous mass showed fairly fibrous capsule and adipocytes with proliferating capillaries.

B) High-power 20X, H&E-stained biopsy of the same patient revealed that adipocytes nuclei are small and bland, compressed at the periphery of cells. Capillaries lined with one layer of bland endothelial cells and containing RBCs.

fairly fibrous capsule, no atypia or mitotic figures (Figure 2). The diagnosis of Dercum’s disease was reached and the patient received symptomatic treatment.

Discussion

The clinical symptoms of Dercum’s disease do not show a specific pattern. Many lipomas can be found in the subcutaneous tissue of the knees, back, neck, thighs, and arms of individuals with this illness. Often, the hands and face are uninvolved. Furthermore, Patients complain of paresthesia in the skin above the lipomas, which is spontaneously painful. A low threshold for pain also seems to be typical for this disease. Normal or glossy skin with dilated superficial veins may also cover lipomas[1,3,6]. Our patient presented with painful swelling in the trunk, upper and lower limbs with sparing head, neck, hands, and feet.

Many other nonspecific symptoms of Dercum’s disease have been reported in the literature. It has a wide range of symptoms such as easy bruising, sleep issues, memory loss, anxiety, depression, and difficulty concentrating; however, these symptoms are not always present. Psychiatric symptoms used to be recognized as crucial symptoms for the diagnostic criteria. These are now recognized as symptoms correlated with adiposis dolorosa, although not all individuals have them, and it is challenging to determine their exact connection to the disease [1]. Our patient didn’t report any psychiatric symptoms. Other reported manifestations in patients with Dercum’s disease include loss of pubic and axillary hair, myxedema, hot flushes, arterial hypertension, early congestive heart failure, cyanosis, dyspnea, and tachypnea. It remains unclear which of these are the major or minor symptoms [3]. Our case his systemic examination was within normal limits.
Dercum disease is categorized into four types based on the afflicted adipose tissue’s location and its relationship with lipomas. Type 1 generalized diffuse: very painful adipose tissue without lipomas. Type 2 generalized nodular: generalized pain in the adipose tissue that is more intense inside and around the lipomas. Type 3 localized nodular: painful adipose tissue exclusively within and around lipomas. Lastly type 4 juxta-articular: deposition of painful solitary fat in the proximity of large joints [1]. This case matched with Dercum’s disease type 2 as he is suffering from painful lipomas.

Laboratory analysis lacks specificity. Acute phase reactant values may be raised because adipose tissue inflammation is the defining feature of Dercum’s disease. Since there are currently no known biomarkers linked to Dercum’s disease, the final diagnosis is being determined based on the results of the histology. The histopathology results of tissue specimens are consistent with those of fatty connective tissues and are identical to lipomas [7], as seen in the histopathology results of our patient (figure 2). Radiological investigations also aid in the detection of subcutaneous nodules [8].

Diagnosis of Dercum’s disease can be challenging for doctors at certain times. Fibromyalgia, which can manifest similarly to Dercum’s disease, is considered a differential diagnosis. However, adiposis dolorosa mostly causes more widespread and intense pain than fibromyalgia. Moreover, familial multiple lipomatosis, which is an autosomal dominant disease, typically manifests as multiple subcutaneous lipomas of varying sizes that are broadly distributed; however, they do not cause incapacitating pain as seen in Derum’s disease [9].

As Dercum’s disease has the potential to develop into a chronic, progressive, and debilitating condition, it is essential to educate patients about the disease state. There are currently no treatments approved by the US Food and Drug Administration for Dercum’s disease, and the effectiveness of treatments that has been attempted is variable. Transdermal lidocaine has been demonstrated to significantly reduce pain by > 60%. The regulation of potential sympathetic nervous system hyperactivity causes pain relief [10]. Patients treated with deoxycholic acid experience a reduction in tumor size, as evidenced by radiographs, as well as a significant improvement in symptoms [11]. A spinal cord stimulator showed a significant improvement in burning pain [12]. Pain in our patient usually relieved by using oral none opioid pain reliever (Acetaminophen).

Primary care physicians and dermatologists must be familiar with Dercum’s disease to ensure that such syndromes are correctly identified in the outpatient setting. Early identification and evaluation are essential to initiate treatment and referral to the appropriate subspecialties, reduce stigmatization, and ultimately improve quality of life. As Dercum’s disease is seen more frequently in obese women [5], our patient, however, had a normal BMI and was a young male patient, making it a unique presentation in the literature.

Conclusions

Diagnosis of Dercum’s disease should be considered in generalized painful lumps. Although the disease is commonly seen in obese women, males with a normal BMI are also affected. In general, the recommended clinical protocol for the evaluation of painful masses includes a comprehensive history and physical examination, serologic markers of acute-phase reactants, and biopsy of suspicious lesions.

Additional Information

Disclosures

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References

Increased Macrophages, and Adipocyte Hypertrophy in Lipedema Thigh Skin and Fat Tissue. J Obes. 2019, 5:10.1155/2019/8747461